

Biomapas is a leading contract research organization specializing in complex clinical research, marketing authorization and post-authorization support in Europe and CIS markets. With more than 16 years of experience and latest geographical expansion, Biomapas is on a way to become a strongest market player.

At the moment Biomapas is looking for **Pharmacovigilance Manager**.

Main Tasks and Responsibilities

- 1. Ensure effective functioning of assigned PV projects in compliance with client agreement, contract; and local and global regulatory guidelines.
- 2. To be responsible for quality management, workload management, compliance management and document management across assigned PV projects.
- 3. Continuously work with internal and external clients to ensure satisfaction.
- 4. Ensure monthly invoices are generated and shared for all assigned PV clients on regular/monthly basis.
- 5. Identify out-of-scope activities and liaise with PV Director and internal departments to initiate and process change orders.
- 6. Provide clarification and/or resolve PV invoicing issues.
- 7. Ensure pharmacovigilance aspects of projects are managed in line with budgets and agreed timelines to achieve client satisfaction.
- 8. Assist with or generates and reviews proposals and costings for pharmacovigilance business.
- 9. Train and mentor PV department staff, as needed. Constantly work towards identification of new training and development opportunities for the PV department.
- 10. Represent PV department during for-cause/maintenance client audits or regulatory authority inspections.
- 11. Represent PV department during pre-qualification client audits.
- 12. Generate and review departmental SOPs and Working Procedures.
- 13. Other responsibilities as assigned by the PV Director

Experience, required skills and competencies

- Life Sciences education.
- More than 5 years of experience in Pharmacovigilance or Project Management field.
- Extensive experience in management of large scale (geographically and content wise) projects.
- Awareness of global regulatory and pharmacovigilance environment.
- Excellent knowledge of English.
- Russian would be considered as an advantage.
- Excellent verbal and written communication, teaming and problem-solving skills.

If you have any questions or would like to find out more about the position, please, contact us by e-mail personalas@biomapas.eu or call directly to HR and Training Manager Raimonda Klimiene +370 698 15736.